

AMENDMENTS TO THE CLAIMS

Claim 1 (original): A syringe and nozzle tip assembly, comprising:

a syringe having a syringe barrel with a front end and a rear end, a piston slidably mounted in the syringe barrel, and a plunger connected to the piston and extending rearwardly through the rear end of the syringe barrel; and

a nozzle tip having a flange, a sleeve frictionally mounted on the front end of the syringe barrel, and a neck with a curved portion and a passage therethrough, wherein the flange includes a recess disposed therein and a filter mounted in the recess.

Claim 2 (original): The syringe and nozzle tip assembly of claim 1, wherein the filter comprises a screen and the screen and curved neck portion are integral with the nozzle tip.

Claim 3 (original): The syringe and nozzle tip assembly of claim 2, wherein the screen has a mesh size of about 105 microns.

Claim 4 (original): The syringe and nozzle tip assembly of claim 1, wherein the outer surface of the syringe barrel is substantially smooth.

Claim 5 (original): The syringe and nozzle tip assembly of claim 1, wherein the recess and filter are configured so that the filter does not contact the end of the syringe barrel.

Claim 6 (original): The syringe and nozzle tip assembly of claim 1, wherein the syringe barrel is transparent.

Claim 7 (original): The syringe and nozzle tip assembly of claim 1, wherein the syringe barrel is made of a material selected from the group consisting of glass and plastic.

Claim 8 (original): The syringe and nozzle tip assembly of claim 1, wherein the nozzle tip and filter are made of low density polyethylene.

Claim 9 (original): The syringe and nozzle tip assembly of claim 1, wherein the nozzle tip is adapted to retain bone regeneration material in the syringe barrel.

Claim 10 (original): The syringe and nozzle tip assembly of claim 1, wherein the flange includes a surface adapted to seat against the syringe barrel when mounted thereon.

Claim 11 (original): The syringe and nozzle tip assembly of claim 1, wherein the nozzle tip is mounted solely by friction fit.

Claim 12 (original): A syringe and nozzle tip assembly, comprising:

a syringe; and

a nozzle tip frictionally mounted on an end of the syringe, the nozzle tip comprising a sleeve, a flange having a surface adapted to seat against the end of the syringe when the nozzle tip

is mounted thereon and a recess disposed therein, a filter mounted in the recess, and a neck having a curved portion and a passage extending therethrough.

Claim 13 (original): The syringe and nozzle tip assembly of claim 12, wherein the nozzle tip is mounted solely by friction fit.

Claim 14 (original): The syringe and nozzle tip assembly of claim 12, wherein the filter is removable.

Claim 15 (original): A syringe and nozzle tip assembly, comprising:

a syringe; and

a nozzle tip mounted on an end of the syringe, the nozzle tip comprising a sleeve, a flange having a surface adapted to seat against the end of the syringe when the nozzle tip is mounted thereon and a recess disposed therein, a filter mounted in the recess, and a neck having a curved portion and a passage extending therethrough, wherein the recess and filter are configured so that the filter does not contact the end of the syringe.

Claim 16 (original): The syringe and nozzle tip assembly of claim 15, wherein the filter is removable.

Claim 17 (original): The syringe and nozzle tip assembly of claim 15, wherein the nozzle tip is mounted frictionally.

mixing the aspirated marrow blood with the bone regeneration material in the syringe until an amount of viscous mass of a mixture of bone regeneration material and marrow blood has formed therein;

removing the nozzle tip from the syringe; and

applying an amount of the viscous mixture to the surgical site.

Claim 22 (new): The method of claim 21, further comprising expelling excess marrow blood in the syringe through the nozzle tip prior to removing the nozzle tip.

Claim 23 (new): A method of using the syringe and nozzle tip assembly of claim 15, comprising:

providing an amount of granular bone regeneration material in the syringe;

aspirating an amount of marrow blood from a surgical site in a patient through the nozzle tip and into syringe;

mixing the aspirated marrow blood with the bone regeneration material in the syringe until an amount of viscous mass of a mixture of bone regeneration material and marrow blood has formed therein;

removing the nozzle tip from the syringe; and

applying an amount of the viscous mixture to the surgical site.

Claim 24 (new): The method of claim 21, further comprising expelling excess marrow blood in the syringe through the nozzle tip prior to removing the nozzle tip.